Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

- 1. (*currently amended*) A method for the production of a coagulant-thrombin from anticoagulated whole blood for formation of a wound healing material, comprising:
 - a) obtaining a volume of anticoagulated whole blood from a subject;
- b) mixing said anticoagulated whole blood with a precipitating agent ethanol at room temperature;
- c) incubating the mixture of b) at room temperature for a time sufficient to produce a cellular and specific plasma component precipitate and a supernatant;
 - d) separating the precipitate from the supernatant; and
- e) recovering the supernatant wherein said supernatant contains a coagulant a thrombin preparation and is in a form suitable for application as a wound healing material comprising 80-90% of prothrombin-thrombin proteins, no detectable fibringen and 20-30% of baseline levels of anti-thrombin III (ATIII).
- 2. (*original*) The method of claim 1, wherein the volume of anticoagulated whole blood is between 8 to 10 ml.
- 3. (previously presented) The method of claim 1, wherein the whole blood is anticoagulated with an anticoagulant selected from the group consisting of acid citrate dextrose (ACD), ACD/mannitol, citrate phosphate dextrose (CPD), and ethylenediaminetetraacetic acid (EDTA).
- 4. (*original*) The method of claim 3, wherein the whole blood is anticoagulated with acid-citrate-dextrose.

Docket No.: 1459.008A

5. (*original*) The method of claim 3, where the whole blood is anticoagulated with ACD/mannitol.

- 6. (*original*) The method of claim 5, wherein the mannitol is present in a concentration of 7.5 mg/ml ACD.
- 7. (*original*) The method of claim 1, wherein the precipitating agent is ethanol.
- 8. (*original*) The method of claim 7, where said ethanol used is at a starting concentration of about 10% to 100%.
- 9. (*original*) The method of claim 8, where said ethanol used is at a starting concentration of about 25% to 95%.
- 10. (*original*) The method of claim 9, where said ethanol used is at a starting concentration of about 50% to 95%.
- 11. (*original*) The method of claim 1, wherein the precipitating agent is a mixture of ethanol and calcium chloride.
- 12. (*original*) The method of claim 1, wherein the incubation step requires less than 45 minutes.
- 13. (*original*) The method of claim 1, wherein the incubation step requires less than 30 minutes.
- 14. (original) The method of claim 1, wherein the coagulant prepared is autologous.
- 15. (original) The method of claim 1, wherein the coagulant prepared is homologous.
- 16. (*original*) The method of claim 1, wherein said separating step is accomplished by centrifuging the mixture.
- 17. (*original*) The method of claim 1, wherein said separating step is accomplished by filtering the mixture.

18. (*original*) The method of claim 1, wherein said separating step is accomplished by a combination of centrifugation and filtration of the mixture.

19-20. (cancelled)

- 21. (currently amended) The method of claim-122, wherein said blood derivative is chosen from the group consisting of a platelet concentrate (PC), platelet rich plasma (PRP), platelet poor plasma (PPP), purified fibrinogen or a mixture thereof to obtain a wound healing composition.
- 22. (currently amended) A method for the production of a coagulant from anticoagulated whole blood for formation of a wound healing material, consisting of:
 - a) obtaining a volume of anticoagulated whole blood from a subject;
- b) mixing said anticoagulated whole blood with a precipitating agentethanol at room temperature;
- c) incubating the mixture of b) at room temperature for a time sufficient to produce a cellular and specific plasma component precipitate and a supernatant;
 - d) separating the precipitate from the supernatant; and
 - e) recovering the supernatant wherein said supernatant contains a coagulant and is in a form suitable for application as a wound healing material thrombin; and
 - f) combining said supernatant with a blood derivative to form a wound healing material.